

Communication

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# A Highly Convergent Total Synthesis of Leustroductsin B.

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Supporting Information Placeholder

**ABSTRACT:** Leustroductsin B exhibits a large variety of biological activities and unique structural features. An efficient and highly convergent total synthesis of Leustroductsin B was achieved in 17 longest linear and 39 total steps by disconnecting the molecule into three fragments having similar levels of complexity. These pieces were connected *via* a highly efficient chelate-controlled addition of a vinyl zincate to an  $\alpha$ -hydroxy ketone and a silicon-mediated cross-coupling. The stereochemistry of the central and western fragments was set catalytically in high yields and excellent *ee* by a zinc-ProPhenol-catalyzed aldol reaction and a palladium-catalyzed asymmetric allylic alkylation.

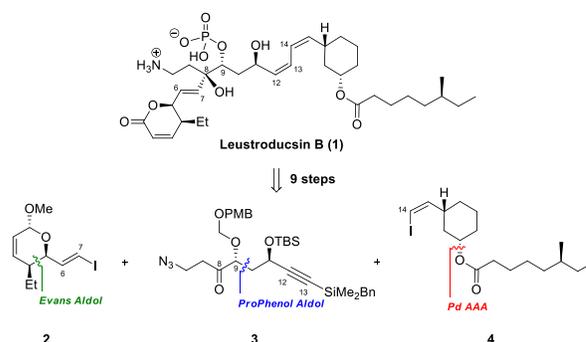
Leustroductsins A-C were first identified and isolated from a culture broth of the soil bacterium *Streptomyces platensis* SANK 60191 in 1993 by Kohama and co-workers at Sankyo.<sup>1</sup> These compounds belong to the phoslactomycin family<sup>2</sup> and exhibit a range of potentially useful bioactivities, such as antibacterial, antifungal, and antitumor activity.<sup>3</sup> They showed high potency and selectivity towards inhibition of protein serine/threonine phosphatase 2A (PP2A), an enzyme that plays a central role in the regulation of cell growth and inhibition of metastasis.<sup>4</sup> Leustroductsin B has specifically shown potent *in vitro* induction of cytokine production by KM-102 cells<sup>3d</sup> as well as an increase of host *in vivo* resistance to *E. Coli* infections<sup>4a</sup> and thrombocytosis induction<sup>3b</sup> when administered to mice. These remarkable biological activities, and its unique highly congested linear structure makes Leustroductsin B a synthetic target of strong interest for the scientific community.<sup>1,5</sup>

Currently, two total syntheses,<sup>6</sup> two formal syntheses,<sup>7</sup> and a semi-synthesis<sup>8</sup> starting from leustroductsin H have been reported. These excellent efforts reveal the difficulty of the challenge as the shortest prior synthesis required 37 linear and 64 total steps, therefore, a more efficient total synthesis of leustroductsin B remains of high interest. As part of our ongoing program aimed at synthesizing highly potent molecules bearing structural complexity in a step and atom economical fashion,<sup>9</sup> we report herein a concise and highly convergent synthesis of leustroductsin B.

Our retrosynthetic approach takes advantage of a highly convergent strategy involving two key disconnections to provide three intermediates with similar size and structural

complexity (Scheme 1). We envisioned that western fragment **2** and central fragment **3** could be coupled *via*  $\alpha$ -alkoxy-directed diastereoselective vinyl zinc addition to generate the C<sub>7</sub>-C<sub>8</sub> bond.<sup>10</sup> Fragment **4** could be united to the rest of the molecule by the modified Hiyama<sup>11</sup> silicon-based palladium-catalyzed cross coupling reaction (C<sub>13</sub>-C<sub>14</sub> bond) developed in our group.<sup>12</sup> To enhance the efficiency of the synthesis, the absolute and relative stereochemistry of fragments **3** and **4** is derived respectively from a zinc-ProPhenol catalyzed aldol reaction<sup>13,14</sup> and a palladium-catalyzed asymmetric allylic alkylation.<sup>15</sup>

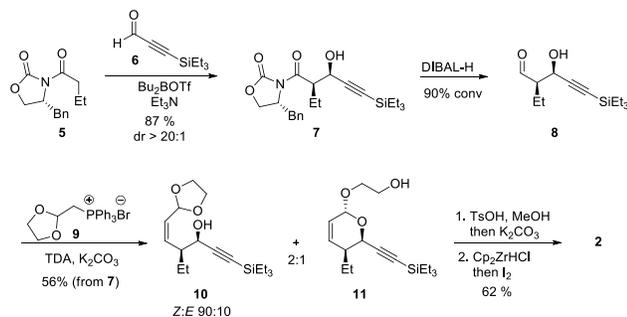
## Scheme 1. Retrosynthetic Analysis of Leustroductsin B.



Vinyl iodide fragment **2** was most efficiently synthesized from commercially available oxazolidinone **5** and propionaldehyde **6** in five steps (Scheme 2). This synthesis is an alternative strategy to the ring-closing metathesis approach recurrently used in the preparation of the 6-membered  $\alpha,\beta$ -unsaturated lactone moiety found in the leustroductsin,<sup>7</sup> phoslactomycin,<sup>16</sup> fostriecyn<sup>17</sup> and cytostatin<sup>18</sup> families. Alcohol **7** was obtained in high yield and excellent diastereomeric ratio following Evan's aldol protocol<sup>19</sup> and the chiral auxiliary was removed using diisobutylaluminum hydride to afford aldehyde **8** as a solution in toluene upon work-up.<sup>20</sup> This aldehyde was treated under mild phase-transfer Wittig reaction conditions in the presence of tris[2-(2-methoxyethoxy)ethyl]amine (TDA) and phosphonium bromide **9** to obtain a mixture of alcohols **10** and **11** in 56% yield over two steps.<sup>21</sup> Ketalization using a catalytic amount of *p*-toluenesulfonic acid in methanol was followed by addition of potassium carbonate to deprotect the terminal alkyne in a *one-pot* procedure. Lastly, hydrozirconation of the terminal alkyne and addition of molecular iodine gave access to key

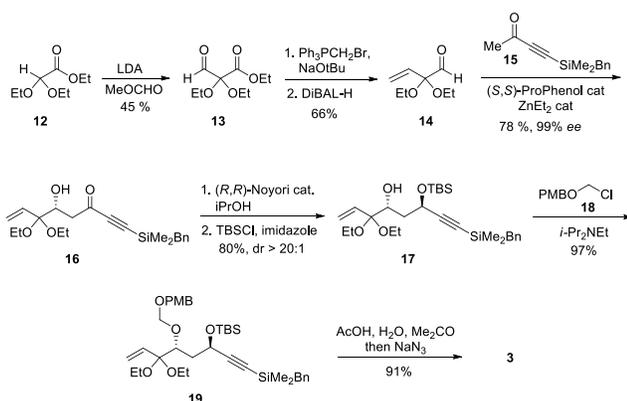
intermediate **2** in 62% yield over two steps on a multigram scale.<sup>22</sup>

### Scheme 2. Synthesis of Eastern Fragment 2.



With fragment **2** in hand, an efficient route to synthesize central key intermediate **3** was developed (Scheme 3). First, carbonyl addition of the enolate of ester **12** to methyl formate followed by acidic hydrolysis (pH 4) of the hemiacetal intermediate delivered aldehyde **13**. The latter was immediately subjected to Wittig olefination conditions followed by reduction of the ester functionality to give aldehyde **14** in good yield. The zinc-ProPhenol catalyzed aldol reaction between aldehyde **14** and ketone **15**<sup>17b</sup> delivered adduct **16** in 78% yield and 99% *ee*.<sup>14</sup> The 1,3-anti relationship of diol **17** was then set by a asymmetric transfer hydrogenation using Noyori's catalyst (*R,R*)-TsDPEN-Ru(*p*-cymene)<sup>23</sup> in 90% yield and > 20:1 *dr*.<sup>17b</sup> Use of achiral reagent Me<sub>4</sub>NB(OAc)<sub>3</sub>H at -35 °C (Evans' conditions)<sup>24</sup> gave moderate 3:1 *dr*. Although the selectivity was lower, the two diastereomers were separable and 1,3-anti diol **17** could be isolated in 61% yield. The undesired diastereomer could also be recycled back to the ynone **16** using manganese dioxide. Protection of the less sterically encumbered hydroxyl group with *tert*-butyldimethylsilylchloride was followed by addition of alkyl chloride **18** to afford **19** in almost quantitative yield. Finally, *one-pot* deprotection of the diethoxyacetal and conversion of the resulting enone into azide **3** was accomplished in 91% yield.

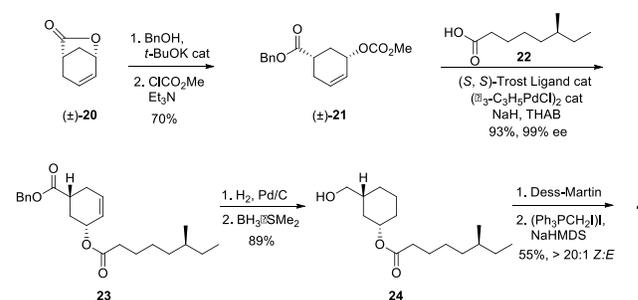
### Scheme 3. Synthesis of Central Fragment 3.



The synthesis of vinyl iodide **4** started from racemic lactone **20** (Scheme 4), which is easily accessible from commercially available 3-cyclohexene-1-carboxylic acid via an iodolactonization/elimination process.<sup>25</sup> Lactone **20** was opened with benzyl alcohol and the resulting hydroxyl group reacted with methylchloroformate to form racemic **21** in 70% yield over two steps. Carbonate **21** was treated with Pd(o), stand-

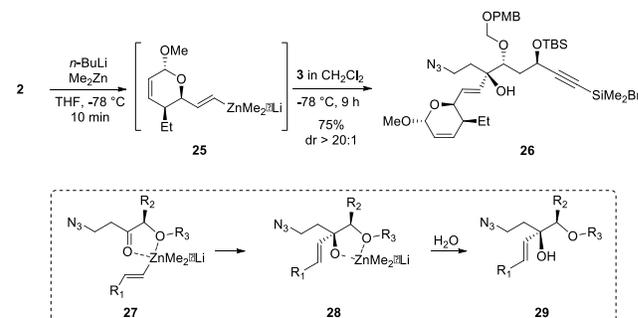
ard (*S,S*)-Trostr ligand and the sodium salt of carboxylic acid **22** (prepared in five synthetic steps from commercially available materials)<sup>26</sup> to deliver ester **23** in 93% yield and 99% *de*.<sup>15</sup> In this particular asymmetric alkylation, the intermediate  $\pi$ -allyl palladium complex is *pseudo-meso*, thus both enantiomers of the starting material can be used in the alkylation. Previous studies showed a so-called memory effect indicating the non-symmetrical nature of the  $\pi$ -allylpalladium intermediate.<sup>27</sup> To insure its full equilibration prior to nucleophilic attack, tetrahexylammonium bromide (THAB) is added to slow the rate of carboxylate attack to maximize the effective symmetry and therefore *ee*. Alcohol **24** was then obtained by hydrogenolysis of ester **23** followed by selective reduction of the carboxylic acid using BH<sub>3</sub>·Me<sub>2</sub>S. Oxidation of alcohol **24** to the corresponding aldehyde was accomplished with Dess-Martin periodinane (DMP) and the resulting aldehyde was subject to a Stork-Zhao olefination to afford vinyl iodide **4** with high *Z*-selectivity.<sup>28</sup>

### Scheme 4. Synthesis of Western Fragment 4.

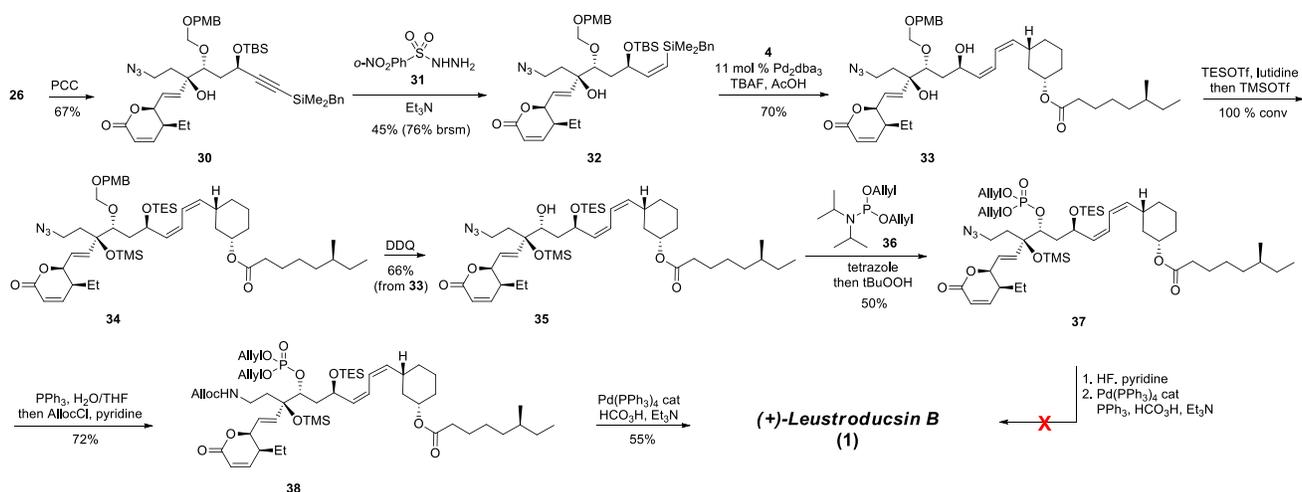


The coupling of the western fragment **2** and central fragment **3** has been the most challenging part of this program (scheme 5). In order to increase the convergent character of our approach for the preparation of leustroducsin B, we investigated the direct addition of vinyl iodide **2** to ketone **3** to generate tertiary allylic alcohol **26** with sensitive functional groups present in the molecule. The desired transformation required formation of metal vinyl species **25** generated from vinyl iodide **2** to add to  $\beta$ -azido ketone **3** in a diastereoselective fashion by chelation to the  $\alpha$ -alkoxy group (**27** to **28**) to finally access alcohol **26**.<sup>10</sup> Extensive screening and reaction optimization<sup>29</sup> revealed that the addition proceeded in high yield and remarkable diastereoselectivity when dimethylzinc<sup>30</sup> was used to generate vinyl zincate complex **25** as the precursor for the chelation-controlled addition. The success of the zincate addition highly depends on the quality of the organometallic reagent employed and the reaction temperature must be kept rigorously under -50 °C in order to avoid decomposition of ketone **3**.

### Scheme 5. Chelation Controlled Addition of 2 to 3.



## Scheme 6. Completion of the Synthesis of Leustroducsin B.



After we successfully coupled fragments **2** and **3**, methyl ketal **26** was oxidized to lactone **30** using pyridinium chlorochromate (PCC). The alkyne moiety was reduced chemoselectively to *cis*-alkene **31** by use of diimide under mild basic conditions<sup>31,32</sup> to avoid significant decomposition of starting material. Cross-coupling between lactone **32** and vinyl iodide **4** was then examined in the presence of a catalytic amount of Pd<sub>2</sub>dba<sub>3</sub>·CHCl<sub>3</sub> and tetrabutylammonium fluoride (TBAF).<sup>12</sup> To our delight, *cis,cis*-diene **33** was isolated in 70% yield. Addition of acetic acid was essential to the success of this transformation as it buffered the reaction mixture.<sup>33</sup> **32** and **33** could easily decompose in presence of residual hydroxide species present in commercial TBAF. The stability of the azide to this series of transformations is particularly noteworthy.

Secondary alcohol **33** was protected with triethylsilyltriflate followed by addition of trimethylsilyltriflate to selectively silylate the hindered tertiary alcohol. The crude material was directly subjected to 1,2-dichloro-4,5-dicyano-1,4-benzoquinone (DDQ) in wet methylene chloride to promote removal of the PMB-ketal functionality and afford secondary alcohol **35** in 66% yield over two steps. Introduction of the phosphate group was accomplished by treatment of alcohol **35** with phosphoramidite reagent **36** in the presence of tetrazole, followed by oxidation of the phosphite intermediate into phosphate **37** in 50% yield.<sup>6</sup> We first attempted to reduce the azide during the final deallylation step using palladium (0) under acidic conditions and an excess of PPh<sub>3</sub>. Although NMR analysis revealed that the allyl groups had been removed, it did not result in the formation of leustroducsin B (**1**). To circumvent this difficulty, a stepwise protocol was pursued. Reduction of azide **37** using standard Staudinger reaction conditions was followed by *in-situ* addition of allylchloroformate and pyridine to deliver carbamate **38** in 72% yield. Similarly to previously reported synthesis of leustroducsin B<sup>6</sup> and to increase the conciseness of our synthesis, we intended to remove both silyl and allyl protecting groups at the last step using an excess of formic acid.<sup>34</sup> Gratifyingly, final deallylation of the phosphonate and the carbamate as well as desilylation of the two hydroxyl groups was successfully achieved using a catalytic amount of Pd(PPh<sub>3</sub>)<sub>4</sub>, formic acid, and triethylamine to reveal deprotected leustroducsin B in 55% yield.<sup>6</sup>

In conclusion, we have reported a highly convergent total synthesis of (+)-leustroducsin B in only 39 total and 17 longest linear steps starting from commercially available material **12**. In comparison to previously reported routes, our concise approach reduced the total number of steps by 40% and exhibits a strong convergent character. The number of longest linear steps has been cut to less than half compared to the previously reported total syntheses of leustroducsin B.<sup>6</sup> Our strategy takes advantage of the preparation of three key intermediates of similar complexity in terms of size and stereochemistry. Particularly noteworthy is the high enantioselectivity of both our zinc-ProPhenol aldol reaction and palladium-catalyzed deracemization using a carboxylate nucleophile. This synthesis also features the exceptionally diastereoselective addition of vinyl zincate **25** to ketone **3** in the presence of sensitive functional groups and a high yielding silicon-based palladium-catalyzed cross-coupling between two highly functionalized molecules. Finally, the last step of the synthesis removed three allyl and two silyl groups all at once, revealing the mildness of these conditions. The brevity and convergent nature of the synthesis reported herein offers an opportunity to explore further the biological effects of chemical modifications in the structure of leustroducsin B.

## ASSOCIATED CONTENT

## Supporting Information

Experiment details, compound characterization data, and spectra. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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## Notes

The authors declare no competing financial interests.

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